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(FILE 'HOME' ENTERED AT 16:50:10 ON 07 AUG 2006)

FILE 'REGISTRY' ENTERED AT 16:50:22 ON 07 AUG 2006

L1 STRUCTURE UPLOADED

L2 8 S L1

L3 164 S L1 SSS FULL

L4 STRUCTURE UPLOADED

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FULL SUBSET SCREEN SEARCH COMPLETED - 164 TO ITERATE

100.0% PROCESSED 164 ITERATIONS 17 ANSWERS

SEARCH TIME: 00.00.01

L5 17 SEA SUB=L3 SSS FUL L4

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L6 147 L3 NOT L5

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FULL ESTIMATED COST ENTRY SESSION 228.14 228.35

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=> s 16

L7 1 L6

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L7 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2004:2877 CAPLUS

DN 140:59667

TI Preparation of 1-[(indol-3-yl)carbonyl]piperazine derivatives for the

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treatment of pain
IN
     Cowley, Phillip Martin; Caulfield, Wilson; Tierney, Jason; Cairns, James;
    Adam-Worrall, Julia; York, Mark
    Akzo Nobel N.V., Neth.
PΑ
     PCT Int. Appl., 38 pp.
so
    CODEN: PIXXD2
DT
    Patent
    English
LA
FAN.CNT 1
     PATENT NO.
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                       KIND
                                                                DATE
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                        A1
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    EP 1549637
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                               20051110
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PRAI EP 2002-77505
                         Α
                               20020621
     WO 2003-EP50226
                         W
                               20030613
os
    MARPAT 140:59667
GI
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AB Title compds. I [R = 1-4 substituents: H, alkyl, halo, etc.; R1 = cyclo(alk(en)yl); R2 = H, Me, ethyl; R3-5, R3'-6' = H, alkyl, alkyloxy, etc.; R6 = H, alkyl, alkyloxy, halo, etc.; R7 = H, alkyl, cycloalkyl, etc.] are prepared For instance, 7-methoxyindole-3-carboxylic acid (preparation

II

given) is alkylated with bromomethylcyclohexane (DMF, NaH), converted to the acid chloride and used to acylate N-ethylpiperazine to give II, isolated as the maleate. Compds. of the invention exhibited activity in a CB1 receptor binding assay and selected compds. significantly increased tail flick latency with an ED50 < 5 μ mol/kg. I are useful in treatment of pain: such as peri-operative pain, chronic pain neuropathic pain, cancer pain and pain and spasticity associated with multiple sclerosis. 639068-09-0P

RL: PAC (Pharmacological activity); PUR (Purification or recovery); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(preparation of 1-[(indol-3-yl)carbonyl]piperazine derivs. for treatment of pain)

RN 639068-09-0 CAPLUS

CN Piperazine, 1-[[1-(1-cyclohexylethyl)-1H-indol-3-yl]carbonyl]-4-ethyl-, (-)- (9CI) (CA INDEX NAME)

Rotation (-).

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
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US
     2006135764 22 JUN 2006
DE 102004056104 24 MAY 2006
        1674464 28 JUN 2006
EΡ
     2006143645 08 JUN 2006
JΡ
WO
     2006058720 08 JUN 2006
GB
        2421183 21 JUN 2006
        2877945 19 MAY 2006
FR
RU
        2276150 10 MAY 2006
CA
        2488034 19 MAY 2006
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FULL ESTIMATED COST 0.44 234.36

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FILE COVERS 1907-1966 FILE LAST UPDATED: 01 May 1997 (19970501/UP)

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